We claim:

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- A vector capable of expressing an α-1,2-mannosidase or a functional part thereof in a methylotrophic yeast strain, comprising a nucleotide sequence coding for said α-1,2-mannosidase or said functional part.
 - $\label{eq:continuous} 2. \ \mbox{The vector of claim 1, wherein said α-1,2-mannosidase is a protein from a fungal species.}$
 - 3. The vector of claim 2, wherein said fungus is Trichoderma reesei.
 - 4. The vector of claim 1, wherein said α -1,2-mannosidase is a protein from a mammalian species.
- 15 5. The vector of claim 4, wherein said α -1,2-mannosidase is murine α -1,2-mannosidase IA or IB.
 - $6. \ \, \text{The vector of claim 1, wherein said α-1,2-mannosidase or said} \\ \text{functional part is tagged with an ER-retention signal.}$
 - $\mbox{7. The vector of claim 6, wherein said ER-retention signal comprises} \label{eq:entropy}$ peptide HDEL.
- 8. The vector of claim 1, wherein the nucleotide sequence coding for 25 said α-1,2-mannosidase or said functional part is operably linked to a promoter and a 3' termination sequence.

- The vector of claim 8, wherein said promoter is the promoter of a gene selected from the group consisting of AOXI, AOXII, GAP, and FLD.
- A vector selected from the group consisting of
 pGAPZMFManHDEL, pGAPZMFManMycHDEL, pPICZBMFManMycHDEL, pGAPZmManHDEL, pGAPZmMycManHDEL, pPIC9mMycManHDEL and pGAPZmMycManHDEL.
- 11. A vector capable of expressing a glucosidase II or a functional part thereof in a methylotrophic yeast strain, comprising a nucleotide sequence coding for said glucosidase II or said functional part.
 - 12. The vector of claim 11, wherein said glucosidase II is a protein from a fungal species.
 - 13. The vector of claim 12, wherein said fungus is Saccharomyces cerevisiae

- ${\bf 14. \ \, The \, vector \, of \, claim \, 11, \, wherein \, said \, glucosidase \, II \, is \, a \, protein}$ ${\bf 20} \qquad from \, a \, mammalian \, species.}$
 - 15. The vector of claim 11, wherein said glucosidase II or said functional part is tagged with an ER-retention signal.
- 25 16. The vector of claim 15, wherein said ER-retention signal comprises peptide HDEL.

- 17. The vector of claim 11, wherein the nucleotide sequence coding for said α -1,2-mannosidase or said functional part is operably linked to a promoter and a 3' termination sequence.
- 5 18. The vector of claim 17, wherein said promoter is the promoter of a gene selected from the group consisting of AOXI, AOXII, GAP, and FLD.
 - 19. A vector having the designation pGAPZAGLSII, pPICZAGLSII, pAOX2ZAGLSII, pYPTIZAGLSII, pGAPADEgIsII, pPICADEgIsII, pAOX2ADEgIsII, pYPTIADEgIsII, pGAPZAgIsIIHDEL and pGAPADEgIsIIHDEL.

- 20. A vector for disrupting the Och1 gene in a methylotrophic yeast strain, comprising a portion of the Och1 gene and a selectable marker gene, wherein said portion of the Och1 gene and said selectable marker gene are linked in such a way to effect the disruption of the genomic Och1 gene in said methylotrophic yeast strain.
 - 21. A vector having the designation pBLURA5'PpOCH1.
- 22. A method of reducing the glycosylation on proteins produced from 20 a methylotrophic yeast, comprising transforming said yeast with any one of the vectors of claims 1-21
 - 23. The method of claim 22, wherein said yeast is Pichia pastoris.
- 25 24. The method of claim 23, wherein said yeast is a *Pichia pastoris* strain selected from GS115 (NRRL Y-15851), GS190 (NRRL Y-18014), PPF1 (NRRL Y-18017), PPY12-OH, yGC4, or derivatives thereof.

- 25. The method of claim 22, 23 or 24, wherein said yeast has been genetically engineered to expresses a heterologous protein.
- 26. A genetically engineered strain of a methylotrophic yeast, wherein

 5 said strain is transformed with at least one of the vectors of claims 1-21.
 - 27. A method of reducing the glycosylation of a heterologous glycoprotein expressed from a methylotrophic yeast, comprising transforming cells of said methylotrophic yeast with at least one of the vectors of claims 1-21, and producing said glycoprotein from the transformed cells.

- 28. A method of producing a glycoprotein with reduced glycosylation in a methylotrophic yeast, comprising transforming cells of said methylotrophic yeast with at least one of the vectors of claims 1-21 and with a nucleotide sequence capable of expressing said glycoprotein in said yeast, and producing said glycoprotein from the transformed cells.
 - 29. A glycoprotein produced by the method of claim 27 or 28.
- 30. The glycoprotein of claim 29, wherein said glycoprotein has a reduced immunogenicity as relative to the glycoprotein produced from a wild type strain of said methylotrophic yeast.
- 31. The glycoprotein of claim 29, wherein said glycoprotein is suitable25 for use in human therapeutics.
 - 32. A kit comprising any of the vectors of claims 1-21.

- 33. The kit of claim 32, further comprising a methylotrophic yeast strain.
 - 34. A kit comprising the methylotrophic yeast strain of claim 26.